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(54) Title: TREATMENT OF DIABETES WITH THIAZOLIDINEDIONE AND SULPHONYLUREA		
(57) Abstract A method for the treatment of diabetes mellitus and conditions associated with diabetes mellitus in a mammal, which method comprises administering an effective non-toxic and pharmaceutically acceptable amount of an insulin sensitiser and a sub-maximal amount of an insulin secretagogue, to a mammal in need thereof; and a pharmaceutical composition for use in such method.		

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TREATMENT OF DIABETES WITH THIAZOLIDINEDIONE AND SULPHONYLUREA

This invention relates to a method of treatment, in particular to a method for the treatment of diabetes mellitus, especially non-insulin dependent diabetes

5 (NIDDM) (or Type 2 diabetes) and conditions associated with diabetes mellitus.

Insulin secretagogues are compounds that promote increased secretion of insulin by the pancreatic beta cells.

The sulphonylureas are well known examples of insulin secretagogues. The sulphonylureas act as hypoglycaemic agents and are used in the treatment of Type 2
10 diabetes. Examples of sulphonylureas include glibenclamide, glipizide, gliclazide, glimepiride, tolazamide and tolbutamide.

European Patent Application, Publication Number 0,306,228 relates to certain thiazolidinedione derivatives disclosed as having hypoglycaemic and hypolipidaemic activity. One particular thiazolidinedione disclosed in EP 0306228 is 5-[4-[2-(N-
15 methyl-N-(2-pyridyl)amino)ethoxy]benzyl]thiazolidine-2,4-dione (hereinafter 'Compound (I)'). WO94/05659 discloses certain salts of Compound (I) including the maleate salt.

Compound (I) is an example of a class of anti-hyperglycaemic agents known as 'insulin sensitisers'. In particular Compound (I) is a thiazolidinedione insulin
20 sensitiser.

European Patent Applications, Publication Numbers: 0008203, 0139421, 0032128, 0428312, 0489663, 0155845, 0257781, 0208420, 0177353, 0319189, 0332331, 0332332, 0528734, 0508740; International Patent Application, Publication
Numbers 92/18501, 93/02079, 93/22445 and United States Patent Numbers 5104888
25 and 5478852, also disclose certain thiazolidinedione insulin sensitisers.

Another series of compounds generally recognised as having insulin sensitiser activity are those typified by the compounds disclosed in International Patent Applications, Publication Numbers WO93/21166 and WO94/01420. These
compounds are herein referred to as 'acyclic insulin sensitisers'. Other examples of
30 acyclic insulin sensitisers are those disclosed in United States Patent Number 5232945 and International Patent Applications, Publication Numbers WO92/03425 and WO91/19702.

Examples of other insulin sensitisers are those disclosed in European Patent Application, Publication Number 0533933, Japanese Patent Application Publication
35 Number 05271204 and United States Patent Number 5264451.

It is now surprisingly indicated that Compound (I) in combination with a sub maximal amount of an insulin secretagogue provides a particularly beneficial effect on glycaemic control, such combination is therefore particularly useful for the treatment

of diabetes mellitus and conditions associated with diabetes. Lowering the dose of the insulin secretagogue in the presence of a full dose of insulin sensitising agent also has the benefit of reducing the likelihood, frequency and/or severity of hypoglycaemic episodes.

5 Accordingly, the invention provides a method for the treatment of diabetes mellitus, especially Type 2 Diabetes, and conditions associated with diabetes in a mammal such as a human, which method comprises administering an effective non-toxic and pharmaceutically acceptable amount of an insulin sensitiser and a sub-maximal amount of an insulin secretagogue, to a mammal in need thereof.

10 In another aspect the invention provides an insulin sensitiser, such as Compound (I), together with a sub-maximal amount of an insulin secretagogue for use in a method for the treatment of diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus.

15 In another aspect the invention provides the use of an insulin sensitiser, such as Compound (I), and a sub-maximal amount of an insulin secretagogue in the manufacture of a composition for the treatment of diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus.

20 It is also considered that the invention encompasses a method for reducing the likelihood, frequency and/or severity of hypoglycaemic episodes, which method comprises administering an effective non-toxic and pharmaceutically acceptable amount of an insulin sensitiser and a sub-maximal amount of an insulin secretagogue.

25 Accordingly, the invention also provides an insulin sensitiser, such as Compound (I), together with an insulin secretagogue for use in reducing the likelihood, frequency and/or severity of hypoglycaemic episodes in the treatment of diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus, wherein the dose of the insulin secretagogue is a sub-maximal dose.

30 In another aspect the invention provides the use of an insulin sensitiser, such as Compound (I), and an insulin secretagogue for the manufacture of a composition for reducing the likelihood, frequency and/or severity of hypoglycaemic episodes in the treatment of diabetes mellitus, especially Type 2 diabetes and conditions associated with diabetes mellitus, wherein the amount of the insulin secretagogue is sub-maximal.

35 The method comprises either co-administration of the insulin sensitiser and the sub-maximal amount of an insulin secretagogue or sequential administration thereof.

Co-administration includes administration of a formulation that includes an insulin sensitiser and a sub-maximal amount of the insulin secretagogue or the essentially simultaneous administration of separate formulations of each agent.

A suitable insulin sensitiser is a thiazolidinedione insulin sensitiser.

5 A suitable thiazolidinedione insulin sensitiser is Compound (I).

Other suitable thiazolidinedione insulin sensitisers include (+) -5-[[4-[(3,4-dihydro-6-hydroxy-2, 5, 7, 8-tetramethyl-2H-1-benzopyran-2-yl)methoxy]phenyl]methyl]-2,4-thiazolidinedione (or troglitazone), 5-[4-[(1-methylcyclohexyl)methoxy]benzyl] thiazolidine-2,4-dione (or ciglitazone), 5-[4-[2-(5-ethylpyridin-2-yl)ethoxy]benzyl] thiazolidine-2,4-dione (or pioglitazone) or 5-[(2-benzyl-2,3-dihydrobenzopyran)-5-ylmethyl]thiazolidine-2,4-dione (or englitazone)

Suitable insulin secretagogues include sulphonylureas.

Suitable sulphonylureas include glibenclamide, glipizide, gliclazide, glimepiride, tolazamide and tolbutamide.

15 Further sulphonylureas include acetohexamide, carbutamide, chlorpropamide, glibornuride, gliquidone, glisentide, glisolamide, glisoxepide, glyclopyamide and glycylamide.

Further suitable insulin secretagogues include repaglinide.

In one particular aspect, the method comprises the administration of 2 to 12 mg of Compound (I), especially when administered per day.

Particularly, the method comprises the administration of 2 to 4, 4 to 8 or 8 to 12 mg of Compound (I) per day.

Particularly, the method comprises the administration of 2 to 4mg of Compound (I), especially when administered per day.

25 Particularly, the method comprises the administration of 4 to 8mg, especially when administered per day.

Particularly, the method comprises the administration of 8 to 12 mg of Compound (I), especially when administered per day.

Preferably, the method comprises the administration of 2 mg of Compound (I), especially when administered per day.

Preferably, the method comprises the administration of 4 mg of Compound (I), especially when administered per day.

Preferably, the method comprises the administration of 8 mg of Compound (I), especially when administered per day.

35 It will be understood that the insulin sensitiser, such as Compound (I) and the insulin secretagogue are each administered in a pharmaceutically acceptable form, including pharmaceutically acceptable derivatives such as pharmaceutically acceptable salts, esters and solvates thereof, as appropriate. In certain instances herein

the names used for the relevant insulin secretagogues may relate to a particular pharmaceutical form of the relevant active agent: It will be understood that all pharmaceutically acceptable forms of the active agent per se are encompassed by this invention, including pharmaceutically acceptable salted forms and pharmaceutically acceptable solvated forms.

Suitable pharmaceutically acceptable salted forms of the insulin sensitisers, such as Compound (I), include those described in the above mentioned patents and patent applications such as in EP 0306228 and WO94/05659 for Compound (I). A preferred pharmaceutically acceptable salt for Compound (I) is a maleate.

Suitable pharmaceutically acceptable solvated forms of the insulin sensitisers, such as Compound (I), include those described in the above mentioned patents and patent applications, such as in EP 0306228 and WO94/05659 for Compound (I), in particular hydrates.

Suitable pharmaceutically acceptable forms of the insulin secretagogue depend upon the particular compound used but include known pharmaceutically acceptable form of the particular compound chosen. Such derivatives are found or are referred to in standard reference texts such as the British and US Pharmacopoeias, Remington's Pharmaceutical Sciences (Mack Publishing Co.), Martindale The Extra Pharmacopoeia (London, The Pharmaceutical Press) (for example see the 31st Edition page 341 and pages cited therein.)

The insulin sensitisers, such as Compound (I) and/or the pharmaceutically acceptable forms thereof, may be prepared using known methods, for example those disclosed in the above mentioned patents and patent applications, such as EP 0306228 and WO94/05659 for Compound (I). The disclosures of the above mentioned patents and patent applications, such as EP 0306228 and WO94/05659, are incorporated herein by reference.

Compound (I) may exist in one of several tautomeric forms, all of which are encompassed by the term Compound (I) as individual tautomeric forms or as mixtures thereof. Compound (I) contains a chiral carbon atom, and hence can exist in up to two stereoisomeric forms, the term Compound (I) encompasses all of these isomeric forms whether as individual isomers or as mixtures of isomers, including racemates.

The insulin secretagogue of choice is prepared according to known methods, such methods are found or are referred to in standard reference texts, such as the British and US Pharmacopoeias, Remington's Pharmaceutical Sciences (Mack Publishing Co.), Martindale The Extra Pharmacopoeia (London, The Pharmaceutical Press) (for example see the 31st Edition page 341 and pages cited therein.)

When used herein the term 'sub-maximal amount' of an insulin secretagogue means an amount lower than (that is less than 100% and typically within the range of

from 5-95% of , for example 75%, 80%, 90% or 95% of) the appropriate non-combination dose for the insulin secretagogue in question, as described or referred to in reference texts such as the British National Formulary (BNF), British and US Pharmacopoeias, Remington's Pharmaceutical Sciences (Mack Publishing Co.),
5 Martindale The Extra Pharmacopoeia (London, The Pharmaceutical Press). For example, for glibenclamide, the maximum dose quoted in the BNF is 15 mg daily; thus a sub-maximal amount of glibenclamide given together with an insulin sensitiser is typically 1.5-12.5 mg daily. For a second example, for gliclazide, the maximum
10 daily dose quoted in the BNF is 320 mg daily; thus a sub-maximal amount of gliclazide given together with an insulin sensitiser is 20-300 mg daily. For a third example, for glipizide, the maximum dose quoted in the BNF is typically 40 mg daily; thus a sub-maximal amount of glipizide given together with an insulin sensitiser is typically 5-30 mg daily. For a fourth example, for tolazamide, the maximum dose
15 quoted in the BNF is 1g daily; thus a sub-maximal amount of tolazamide given together with an insulin sensitiser is typically 50-950 mg daily. For a fifth example, for tolbutamide, the maximum dose quoted in the BNF is 2g daily; thus a sub-maximal amount of tolbutamide given together with an insulin sensitiser is typically 100 mg to 1.75g daily.

When used herein the term 'conditions associated with diabetes' includes those
20 conditions associated with diabetes mellitus itself and complications associated with diabetes mellitus.

'Conditions associated with diabetes mellitus itself' include hyperglycaemia, insulin resistance, including acquired insulin resistance. Further conditions associated with diabetes mellitus itself include hypertension and cardiovascular disease,
25 especially atherosclerosis and conditions associated with insulin resistance. Conditions associated with insulin resistance include polycystic ovarian syndrome and steroid induced insulin resistance and gestational diabetes.

'Complications associated with diabetes mellitus' includes renal disease, especially renal disease associated with Type 2 diabetes, neuropathy and retinopathy.

30 Renal diseases associated with Type 2 diabetes include nephropathy, glomerulonephritis, glomerular sclerosis, hypertensive nephrosclerosis and end stage renal disease. Additional renal diseases associated with Type 2 diabetes include nephrotic syndrome.

As used herein the term 'pharmaceutically acceptable' embraces both human
35 and veterinary use: for example the term 'pharmaceutically acceptable' embraces a veterinarily acceptable compound.

For the avoidance of doubt, when reference is made herein to scalar amounts, including mg amounts, of Compound (I) in a pharmaceutically acceptable form, the

scalar amount referred to is made in respect of Compound (I) *per se*: For example 2 mg of Compound (I) in the form of the maleate salt is that amount of maleate salt which contains 2 mg of Compound (I).

Diabetes mellitus is preferably Type 2 diabetes.

5 Suitably the insulin sensitiser is the first administered agent.

In the present treatment the insulin sensitiser is administered at its normal, appropriate dose, for example Compound (I) is administered at a dose selected from 2-12mg per day, for example 1, 2, 4 or 8 mg per day.

Glycaemic control as referred to herein may be characterised using
10 conventional methods, for example by measurement of a typically used index of glycaemic control such as fasting plasma glucose or glycosylated haemoglobin (HbA1c). Such indices are determined using standard methodology, for example those described in: Tiescher A, Richterich and P., Schweiz. med. Wschr. 101 (1971), 345 and 390 and Frank P., 'Monitoring the Diabetic Patient with Glycosolated
15 Hemoglobin Measurements', Clinical Products 1988.

In the method of the invention, the active medicaments are preferably administered in pharmaceutical composition form. As indicated above, such compositions can include both medicaments or one only of the medicaments.

Accordingly, in one aspect the present invention also provides a
20 pharmaceutical composition comprising an insulin sensitiser, such as Compound (I) and especially 2 to 12 mg thereof, a sub-maximal amount of an insulin secretagogue and a pharmaceutically acceptable carrier therefor.

Such compositions may be prepared by admixing an insulin sensitiser, such as Compound (I) and especially 2 to 12 mg thereof, a sub-maximal amount of an
25 insulin secretagogue and a pharmaceutically acceptable carrier therefor.

Usually the compositions are adapted for oral administration. However, they may be adapted for other modes of administration, for example parenteral administration, sublingual or transdermal administration.

The compositions may be in the form of tablets, capsules, powders, granules,
30 lozenges, suppositories, reconstitutable powders, or liquid preparations, such as oral or sterile parenteral solutions or suspensions.

In order to obtain consistency of administration it is preferred that a composition of the invention is in the form of a unit dose.

Unit dose presentation forms for oral administration may be tablets and
35 capsules and may contain conventional excipients such as binding agents, for example syrup, acacia, gelatin, sorbitol, tragacanth, or polyvinylpyrrolidone; fillers, for example lactose, sugar, maize-starch, calcium phosphate, sorbitol or glycine; tableting lubricants, for example magnesium stearate; disintegrants, for example

starch, polyvinylpyrrolidone, sodium starch glycollate or microcrystalline cellulose; or pharmaceutically acceptable wetting agents such as sodium lauryl sulphate.

The compositions are preferably in a unit dosage form in an amount appropriate for the relevant daily dosage.

5 Suitable dosages for the insulin sensitisers include those disclosed in the abovementioned patents and patent applications.

Suitable dosages, including unit dosages, of Compound (I) comprise 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11 or 12 mg of Compound (I).

10 Particular dosages of Compound (I) are 2mg/day, 4mg/day, including 2mg twice per day, and 8 mg/day, including 4mg twice per day.

In the treatment the medicaments may be administered from 1 to 6 times a day, but most preferably 1 or 2 times per day.

15 The solid oral compositions may be prepared by conventional methods of blending, filling or tableting. Repeated blending operations may be used to distribute the active agent throughout those compositions employing large quantities of fillers. Such operations are of course conventional in the art. The tablets may be coated according to methods well known in normal pharmaceutical practice, in particular with an enteric coating.

20 Oral liquid preparations may be in the form of, for example, emulsions, syrups, or elixirs, or may be presented as a dry product for reconstitution with water or other suitable vehicle before use. Such liquid preparations may contain conventional additives such as suspending agents, for example sorbitol, syrup, methyl cellulose, gelatin, hydroxyethylcellulose, carboxymethylcellulose, aluminium stearate gel, hydrogenated edible fats; emulsifying agents, for example lecithin, sorbitan
25 monooleate, or acacia; non-aqueous vehicles (which may include edible oils), for example almond oil, fractionated coconut oil, oily esters such as esters of glycerine, propylene glycol, or ethyl alcohol; preservatives, for example methyl or propyl p-hydroxybenzoate or sorbic acid; and if desired conventional flavouring or colouring agents.

30 For parenteral administration, fluid unit dosage forms are prepared utilizing the compound and a sterile vehicle, and, depending on the concentration used, can be either suspended or dissolved in the vehicle. In preparing solutions the compound can be dissolved in water for injection and filter sterilized before filling into a suitable vial or ampoule and sealing. Advantageously, adjuvants such as a local anaesthetic, a
35 preservative and buffering agent can be dissolved in the vehicle. To enhance the stability, the composition can be frozen after filling into the vial and the water removed under vacuum. Parenteral suspensions are prepared in substantially the same manner, except that the Compound (I) is suspended in the vehicle instead of being

dissolved, and sterilization cannot be accomplished by filtration. The compound can be sterilized by exposure to ethylene oxide before suspending in the sterile vehicle. Advantageously, a surfactant or wetting agent is included in the composition to facilitate uniform distribution of the compound.

5 Compositions may contain from 0.1% to 99% by weight, preferably from 10-60% by weight, of the active material, depending upon the method of administration.

 Composition may, if desired, be in the form of a pack accompanied by written or printed instructions for use.

10 The compositions are prepared and formulated according to conventional methods, such as those disclosed in standard reference texts, for example the British and US Pharmacopoeias, Remington's Pharmaceutical Sciences (Mack Publishing Co.), Martindale The Extra Pharmacopoeia (London, The Pharmaceutical Press) and Harry's Cosmeticology (Leonard Hill Books) (for example see the 31st Edition page
15 341 and pages cited therein.)

 The present invention also provides a pharmaceutical composition comprising an insulin sensitiser, such as Compound (I) and especially 2 to 12 mg thereof, a sub-maximal amount of an insulin secretagogue and a pharmaceutically acceptable carrier therefor, for use as an active therapeutic substance.

20 The invention also provides the use of an insulin sensitiser, such as Compound (I) and especially 2 to 12 mg thereof, a sub-maximal amount of an insulin secretagogue for the manufacture of a medicament for the treatment of diabetes mellitus and conditions associated with diabetes.

 In particular, the present invention provides a pharmaceutical composition
25 comprising an insulin sensitiser, such as Compound (I) and especially 2 to 12 mg thereof, a sub-maximal amount of an insulin secretagogue and a pharmaceutically acceptable carrier therefor, for use in the treatment of diabetes and conditions associated with diabetes.

 A range of 2 to 4mg includes a range of 2.1 to 4, 2.2 to 4, 2.3 to 4, 2.4 to 4,
30 2.5 to 4, 2.6 to 4, 2.7 to 4, 2.8 to 4, 2.9 to 4 or 3 to 4mg.

 A range of 4 to 8mg includes a range of 4.1 to 8, 4.2 to 8, 4.3 to 8, 4.4 to 8, 4.5 to 8, 4.6 to 8, 4.7 to 8, 4.8 to 8, 4.9 to 8, 5 to 8, 6 to 8 or 7 to 8mg.

 A range of 8 to 12 mg includes a range of 8.1 to 12, 8.2 to 12, 8.3 to 12, 8.4 to 12, 8.5 to 12, 8.6 to 12, 8.7 to 12, 8.8 to 12, 8.9 to 12, 9 to 12, 10 to 12 or 11 to
35 12mg.

 No adverse toxicological effects are expected for the compositions or methods of the invention in the above mentioned dosage ranges.

Compositions of Compound (I)

Preparation of Concentrate: Tabletting concentrate was prepared using the following materials

Ingredient	Quantity (%)
Milled Compound (I) as maleate salt	13.25 (pure maleate salt)
Sodium Starch Glycollate	5.00
Hydroxypropyl Methylcellulose 2910	5.00
Microcrystalline Cellulose (Avicel PH102)	20.0
Lactose Monohydrate, regular grade	to 100
Purified water	*

* Removed during processing.

The concentrate was then formulated into tablets using the following:

Tablet Strength	Quantity (mg per Tablet)			
	1.0mg	2.0mg	4.0mg	8.0mg
Active Ingredient:				
Compound (I) maleate Concentrate granules	10.00	20.00	40.00	80.00
Other Ingredients:				
Sodium Starch Glycollate	6.96	6.46	5.46	10.92
Microcrystalline Cellulose (Avicel PH102)	27.85	25.85	21.85	43.70
Lactose monohydrate, (Pharmatose DCL15),	104.44	96.94	81.94	163.88
Magnesium Stearate	0.75	0.75	0.75	1.50
Total Weight of Tablet Core	150.0	150.0	150.0	300.0
Opadry	4.5	4.5	4.5	9.0
Total Weight of Film Coated Tablet	154.5	154.5	154.5	309.0

Compositions for other active agents are as described in the above mentioned publications.

Claims

1. A method for the treatment of diabetes mellitus and conditions associated with diabetes mellitus in a mammal, which method comprises administering an
5 effective non-toxic and pharmaceutically acceptable amount of an insulin sensitiser and a sub-maximal amount of an insulin secretagogue, to a mammal in need thereof.
2. A method according to claim 1, wherein the insulin secretagogue is a
10 sulphonylurea.
3. A method according to claim 1, wherein the insulin secretagogue is glibenclamide, glipizide, gliclazide, glimepiride, tolazamide, tolbutamide, acetohexamide, carbutamide, chlorpropamide, glibornuride, gliquidone, glisentide, glisolamide,
15 glisoxepide, glycopyramide and glycyllamide or repaglinide.
4. A method according to any one of claims 1 to 3, wherein the insulin sensitiser is 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl]thiazolidine-2,4-dione (Compound I) or a tautomeric form thereof and/or a pharmaceutically acceptable
20 derivative thereof.
5. A method according to any one of claims 1 to 4, which comprises the administration of 2 to 12 mg of Compound (I).
- 25 6. A method according to any one of claims 1 to 5, which comprises the administration of 2 to 4, 4 to 8 or 8 to 12 mg of Compound (I).
7. A method according to any one of claims 1 to 6, which comprises the administration of 2 to 4mg of Compound (I).
30
8. A method according to any one of claims 1 to 6, which comprises the method the administration of 4 to 8mg of Compound (I).
9. A method according to any one of claims 1 to 6, which comprises the
35 administration of 8 to 12 mg of Compound (I).

10. A method according to any one of claims 1 to 6, which comprises the administration of 2 mg of Compound (I), or a tautomeric form thereof and/or a pharmaceutically acceptable derivative thereof.
- 5 11. A method according to any one of claims 1 to 6, which comprises the administration of 4 mg of Compound (I).
12. A method according to any one of claims 1 to 6, which comprises the administration of 8 mg of Compound (I).
- 10 13. A method according to claim 1, wherein the insulin sensitiser is (+) -5-[[4-[(3,4-dihydro-6-hydroxy-2, 5, 7, 8-tetramethyl-2H-1-benzopyran-2-yl)methoxy]phenyl]methyl]-2,4-thiazolidinedione (or troglitazone), 5-[4-[(1-methylcyclohexyl)methoxy]benzyl] thiazolidine-2,4-dione (or ciglitazone), 5-[4-
15 [2-(5-ethylpyridin-2-yl)ethoxy]benzyl] thiazolidine-2,4-dione (or pioglitazone) or 5-[(2-benzyl-2,3-dihydrobenzopyran)-5-ylmethyl]thiazolidine-2,4-dione (or englitazone); or a tautomeric form thereof and/or a pharmaceutically acceptable derivative thereof.
- 20 14. A pharmaceutical composition comprising an insulin sensitiser, a sub-maximal amount of an insulin secretagogue and a pharmaceutically acceptable carrier therefor.
- 25 15. A composition according to claim 14, wherein the insulin secretagogue is a sulphonylurea.
16. A composition according to claim 14 or claim 15, wherein the insulin secretagogue is glibenclamide, glipizide, gliclazide, glimepiride, tolazamide or tolbutamide, acetohexamide, carbutamide, chlorpropamide, glibornuride,
30 gliquidone, glisentide, glisolamide, glisoxepide, glyclopyamide and glycyclamide or repaglinide.
17. A composition according to any one of claims 14 to 16, wherein the insulin sensitiser is Compound (I)
- 35 18. A composition according to any one of claims 14 to 17, which comprises 2 to 12 mg of Compound (I).

19. A pharmaceutical composition comprising an insulin sensitiser, a sub-maximal amount of an insulin secretagogue and a pharmaceutically acceptable carrier therefor, for use as an active therapeutic substance.
- 5 20. A pharmaceutical composition comprising an insulin sensitiser, a sub-maximal amount of an insulin secretagogue and a pharmaceutically acceptable carrier therefor, for use in the treatment of diabetes mellitus and conditions associated with diabetes mellitus.
- 10 21. A composition according to any one of claims 14, 19 or 20, wherein the insulin sensitiser is (+) -5-[[4-[(3,4-dihydro-6-hydroxy-2, 5, 7, 8-tetramethyl-2H-1-benzopyran-2-yl)methoxy]phenyl]methyl]-2,4-thiazolidinedione (or troglitazone), 5-[4-[(1-methylcyclohexyl)methoxy]benzyl] thiazolidine-2,4-dione (or ciglitazone), 5-[4-[2-(5-ethylpyridin-2-yl)ethoxy]benzyl] thiazolidine-2,4-dione (or pioglitazone) or 5-[(2-benzyl-2,3-dihydrobenzopyran)-5-ylmethyl]thiazolidine-2,4-dione (or englitazone); or a tautomeric form thereof
15 and/or a pharmaceutically acceptable derivative thereof.

INTERNATIONAL SEARCH REPORT

In. tional Application No

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A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 A61K31/64 A61K31/44 //(A61K31/64,31:44)

According to International Patent Classification(IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	KELLEY D.E. ET AL: "Troglitazone" CURRENT OPINION IN ENDOCRINOLOGY AND DIABETES, 1998, 5/2 (90-96), XP002080820 United States see page 93, column 2 ---	1-4, 13-16, 19-21
X	SCHEEN AJ ET AL: "Oral antidiabetic agents. A guide to selection." DRUGS, FEB 1998, 55 (2) P225-36, XP002080821 NEW ZEALAND see page 232, column 2, paragraph 1; figure 1 --- -/--	1-4, 13-16, 19-21

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

16 October 1998

Date of mailing of the international search report

02/11/1998

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INTERNATIONAL SEARCH REPORT

In .tional Application No

PCT/GB 98/02109

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	IWAMOTO Y ET AL: "EFFECT OF COMBINATION THERAPY OF TROGLITAZONE AND SULPHONYLUREAS INPATIENTS WITH TYPE 2 DIABETES WHO WERE POORLY CONTROLLED BY SULPHONYLUREA THERAPY ALONE" DIABETIC MEDICINE, vol. 13, no. 4, April 1996, pages 365-370, XP002064289 see abstract ---	1-4, 13-16, 19-21
E	WO 98 36755 A (WARNER LAMBERT CO ;WHITCOMB RANDALL W (US)) 27 August 1998 see page 4, line 30; table 5 see page 71, paragraph 2; claims 1-8 ---	1-21
E	EP 0 861 666 A (TAKEDA CHEMICAL INDUSTRIES LTD) 2 September 1998 see page 8, line 32-42 see page 9, line 45-55 see page 13; claims 24-26; example 2 ---	1-21
A	SCHEEN AJ: "Drug treatment of non-insulin-dependent diabetes mellitus in the 1990s. Achievements and future developments." DRUGS, SEP 1997, 54 (3) P355-68, XP002080822 NEW ZEALAND see page 359; figures 1,2 see page 363, column 1, paragraph 2 ---	1-21
A	WO 93 03724 A (UPJOHN CO) 4 March 1993 see claims 3,4,10,11 ---	1-21
X	EP 0 749 751 A (TAKEDA CHEMICAL INDUSTRIES LTD) 27 December 1996 see page 9, line 42-55; claims 24-26; example 2 see page 8, line 32-34 -----	1-21

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/GB 98/02109

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9836755	A	27-08-1998	NONE
EP 0861666	A	02-09-1998	AU 5603496 A 09-01-1997 CA 2179584 A 21-12-1996 CN 1145783 A 26-03-1997 CZ 9601811 A 15-01-1997 EP 0749751 A 27-12-1996 HU 9601698 A 28-05-1997 JP 9067271 A 11-03-1997 JP 10167986 A 23-06-1998 NO 962606 A 23-12-1996 SK 79496 A 08-01-1997
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INTERNATIONAL SEARCH REPORT

International application No.

PCT/GB 98/02109

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Although claim(s) 1-13
is(are) directed to a method of treatment of the human/animal
body, the search has been carried out and based on the alleged
effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such
an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all
searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment
of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report
covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is
restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.